

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)	Application Number		10581413	
	Filing Date		2006-06-23	
	First Named Inventor	Shilara, Kenya		
	Art Unit	1644		
	Examiner Name	Not Yet Assigned		
Attorney Docket Number		00005.001295		

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	3	2002-539079	JP		2002-11-19	Millennium Pharmaceuticals Incorporated		<input type="checkbox"/>

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4	WO 2002/010743	WO		2002-02-07	Ortho-Mcneil Pharmaceutical, Inc.		<input type="checkbox"/>
5	2002-544173	JP		2002-12-24	Immunomedics Inc.		<input type="checkbox"/>
6	WO 2002/012347	WO		2002-02-14	Immunomedics Inc.		<input type="checkbox"/>
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	1	GOLDENBERG, "Trastuzumab, a Recombinant DNA-Derived Humanized Monoclonal Antibody, a Novel Agent for the Treatment of Metastatic Breast Cancer", Clinical Therapeutics, Vol. 21, No. 2 (1999), pages 309-317	<input type="checkbox"/>
	2	CZUCZMAN, ET AL, "Treatment of Patients With Low-Grade B-Cell Lymphoma With the Combinatio of Chimeric Anti-CD20 Monoclonal Antibody and CHOP Chemotherapy", Journal of Clinical Oncology, Vol. 17, No. 1 (1999), pages 268-276	<input type="checkbox"/>
	3	FRIEDBERG, ET AL, "Combination Immunotherapy with Rituximab and Interleukin 2 in Patients with Relapsed or Refractory Follicular Non-Hodgkin's Lymphoma", British Journal of Hematology, Vol. 117 (2002), pages 828-834	<input type="checkbox"/>
	4	VAN DER KOLK, LE, et al., "Treatment of Relapsed B-Cell non-Hodgkin's Lymphoma With a Combinatio of Chimeric Anti-CD20 Monoclonal Antibodies (rituximab) and G-CSF: Final Report on Safety and Efficacy", Leukemia, Vol. 17 (2003), pages 1658-1664	<input type="checkbox"/>
	5	TAKEUCHI, "Chemical Therapy for Adult ALL", Igaku no Ayumi (Progress Med. Sci.), Vol. 190, No. 5 (1999), pages 474-480	<input type="checkbox"/>

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6	Ohnishi, "Standard Therapy of CML", Igaku no Ayumi (Progress Med. Sci.), Vol. 190, No. 5 (1999), pages 481-485	<input type="checkbox"/>
7	KUSUMOTO, "Usage of cytokine at therapy of acute leukemia", Igaku no Ayumi (Progress Med. Sci.), Vol. 190, No. 5 (1999), pages 522-529	<input type="checkbox"/>

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- ☐ That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.97(e)(1).

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- ☐ See attached certification statement.
- ☐ Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- ☒ None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Lawrence S. Perry/	Date (YYYY-MM-DD)	2007-05-17
Name/Print	Lawrence S. Perry	Registration Number	31865

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